

WHAT IS CLAIMED IS:

1 1. A substantially purified immunoglobulin
2 polypeptide that specifically binds to a human type beta
3 platelet-derived growth factor receptor (β PDGF-R), wherein
4 binding of the polypeptide has one or more of the following
5 effects:

6 i) inhibition of PDGF BB or AB binding to the
7 β PDGF-R;
8 ii) inhibition of PDGF-induced β PDGF-R
9 phosphorylation;
10 iii) inhibition of PDGF-induced dimerization of
11 β PDGF-R;
12 iv) inhibition of PDGF-induced mitogenesis of
13 cells displaying human β PDGF-R; and
14 v) inhibition of PDGF-induced chemotaxis and
15 migration of cells displaying β PDGF-R.

1 2. An immunoglobulin polypeptide of claim 1, wherein
2 the polypeptide is a monoclonal antibody.

1 3. An immunoglobulin polypeptide of claim 2, wherein
2 the monoclonal antibody is antibody 2A1E2.

1 4. A substantially purified polypeptide having an
2 amino acid sequence substantially identical to a sequence of a
3 complementarity determining region of an immunoglobulin
4 polypeptide of claim 1.

1 5. An immunoglobulin polypeptide of claim 1, wherein
2 the polypeptide is linked to a detectable label.

1 6. An immunoglobulin polypeptide of claim 1, wherein
2 the polypeptide is chimeric.

1 7. A substantially purified immunoglobulin
2 polypeptide that specifically recognizes an epitope which lies

3 in the second Ig-like domain in the extracellular region of the
4 β PDGF-R.

1 8. A composition comprising a monoclonal antibody or
2 binding fragment thereof that binds to the human β PDGF-R, which
3 antibody or fragment inhibits in vivo binding of PDGF BB or AB
4 to the receptor.

1 9. An isolated nucleic acid having a sequence
2 substantially identical to a nucleic acid coding for an
3 immunoglobulin polypeptide or a binding fragment thereof,
4 wherein binding of the polypeptide or fragment to a human
5 β PDGF-R has one or more of the following effects:

6 i) inhibition of PDGF BB or AB binding to the
7 β PDGF-R;

8 ii) inhibition of PDGF-induced β PDGF-R
9 phosphorylation;

10 iii) inhibition of PDGF-induced dimerization of
11 β PDGF-R;

12 iv) inhibition of PDGF-induced mitogenesis of
13 cells displaying the human β PDGF-R; and

14 v) inhibition of PDGF-induced chemotaxis and
15 migration of cells displaying β PDGR-R.

1 10. A nucleic acid of claim 9, wherein the nucleic
2 acid is operably linked to a promoter.

1 11. A nucleic acid of claim 10, wherein the promoter
2 and the nucleic acid are contained in an expression vector.

1 12. A cell line transfected, transformed, or infected
2 with a nucleic acid of claim 9.

1 13. A method of producing a substantially purified
2 immunoglobulin polypeptide, or binding fragment thereof, which
3 binds to a human type beta PDGF receptor (β PDGF-R), wherein the
4 binding of the polypeptide or fragment to the β PDGF-R has one

5 or more of the following effects: inhibition of PDGF BB or AB
6 binding to the β PDGF-R; inhibition of PDGF-induced β PDGF-R
7 phosphorylation; inhibition of PDGF-induced dimerization of the
8 β PDGF-R; and inhibition of PDGF-induced mitogenesis of cells
9 displaying human β PDGF-R; and inhibition of PDGF-induced
10 chemotaxis and migration of cells displaying human β -PDGF; the
11 method comprising:

12 i) growing a cell line comprising a nucleic acid
13 encoding the immunoglobulin polypeptide; and
14 ii) harvesting the immunoglobulin polypeptide.

1 14. A method of claim 13, wherein the cell line is a
2 hybridoma.

1 15. A method of claim 14, wherein the hybridoma is
2 ATCC no. HB10938.

1 16. A method of claim 13, wherein the immunoglobulin
2 polypeptide is a monoclonal antibody.

1 17. A method of treating a human having a
2 PDGF-mediated disease involving proliferation, migration or
3 chemotaxis of smooth muscle cells, comprising administering to
4 the patient a therapeutically effective dose of at least one
5 immunoglobulin polypeptide according to claim 1, or fragments
6 of the immunoglobulin polypeptide, and a pharmaceutically
7 acceptable carrier.

1 18. An isolated cell line designated as ATCC no.
2 HB10938.

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